AFRIN NODRIP ALLERGY SINUS NIGHT- oxymetazoline hydrochloride spray Bayer HealthCare LLC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Afrin NoDrip Allergy Sinus Night UI1612461

Drug Facts

Active ingredient Purpose

Oxymetazoline hydrochloride 0.05%......Nasal decongestant

Uses

- temporarily relieves nasal congestion due to:
- common cold •hay fever
- upper respiratory allergies
- reduces swelling of nasal passages so you can breathe more freely
- temporarily relieves sinus congestion and pressure

Warnings

- Ask a doctor before use if you have
- heart disease high blood pressure
- thyroid disease diabetes
- trouble urinating due to an enlarged prostate gland

When using this product

- do not use more than directed
- do not use for more than 3 days. Use only as directed.

Frequent or prolonged use may cause nasal congestion to

recur or worsen.

- temporary discomfort such as burning, stinging, sneezing, or an increase in nasal discharge may occur
- use of this container by more than one person may spread infection

Stop use and ask a doctor if symptoms persist

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• adults and children 6 to under 12 years of age (with adult supervision): 2 or 3 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour

period.

• children under 6 years of age: ask a doctor

To Use: Shake well before use. Hold white tabs, press grooved area of cap firmly and turn counter clockwise. Before using the first time, prime metered pump by depressing pump firmly several times. To spray, hold bottle with thumb at base and nozzle between first and second fingers. Without tilting head, insert nozzle into nostril. Fully depress rim with a firm, even stroke and sniff deeply. Wipe nozzle clean after use. Secure cap after use.

Other information

- store between 20° to 25°C (68° to 77°F)
- retain carton for future reference on full labeling

Inactive ingredients benzalkonium chloride solution, benzyl alcohol, dibasic sodium phosphate, edetate disodium, flavor, glycerin, microcrystalline cellulose and carboxymethylcellulose sodium, monobasic sodium phosphate, polyethylene glycol, povidone, purified water

Questions or comments? 1-800-317-2165

Carton label 15 mL

Oxymetazoline HCl

Nasal Solution-Nasal Decongestant

AFRIN

NODRIP

Wont drive from nose or down throat

Allergy Sinus

Night

with soothing chamomile scent

PUMP MIST

Instant Congestion

Relief from Allergies

for a More Restful Night

Reduces

Swelling of

Nasal

Passages

1/2 FL OZ (15mL)



AFRIN NODRIP ALLERGY SINUS NIGHT

oxymetazoline hydrochloride spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11523-0034
Route of Administration	NASAL		
Active Ingredient/Active N			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE -	OXYMETAZOLINE	0.5 mg

UNII:8 VLN5B44ZY)	HYDROCHLORIDE	in 1 mL
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Inactive Ingredients		
Ingredient Name	Strength	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679 OBS 311)		
SODIUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2SW)		
WATER (UNII: 059QF0KO0R)		
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)		
GLYCERIN (UNII: PDC6A3C0OX)		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
PO VIDO NE (UNII: FZ989 GH94E)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)		
HIBISCUS BIFURCATUS WHOLE (UNII: 60F5JKG79P)		

l	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:11523-0034-1	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/09/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	11/09/2020	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 11/2020 Bayer HealthCare LLC.